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(71) Applicant: **THE DOW CHEMICAL COMPANY**
2030 Dow Center Abbott Road P.O. Box 1967
Midland, MI 48640(US)

(72) Inventor: **Hefner, Robert E., Jr.**
109 Wedgewood
Lake Jackson Texas 77566(US)

(72) Inventor: **Haynes, Deborah I.**
1610 W. 10th.
Freeport Texas 77541(US)

(74) Representative: **Sternagel, Hans-Günther, Dr. et al,**
Patentanwälte Dr. Michael Hann Dr. H.-G. Sternagel
Sander Aue 30
D-5060 Bergisch Gladbach 2(DE)

(54) Improved rheology low class, high impact strength powder coating resins.

(57) Novel advanced epoxy resins are disclosed which are the product resulting from (I) polymerizing in the presence of a catalytic quantity of a suitable polymerization catalyst (A) the reaction product of (1) a mixture consisting of (a) a diglycidyl ether of a dihydric phenol such as a diglycidyl ether of bisphenol A and (b) a diglycidyl ether of a mono-, di- or polyetherdiol such as a diglycidyl ether of a polypropylene glycol with (2) a compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group such as methacrylic acid; with (B) a monomer mixture containing (1) at least one vinyl aromatic monomer such as styrene, (2) a compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group such as methacrylic acid and optionally (3) a hydroxyalkyl acrylate or methacrylate or an alkyl acrylate or methacrylate such as hydroxyethylacrylate; and (II) advancing in the presence of a catalytic quantity of a suitable advancement catalyst, the product from (I) with (C) a dihydric phenol such as bisphenol A.

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IMPROVED RHEOLOGY LOW GLOSS,
HIGH IMPACT STRENGTH POWDER COATING RESINS

Advanced epoxy resin compositions which have been modified by the in situ formation of the copolymerization product of a vinyl aromatic monomer and minor amounts of a compound containing a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group are taught by Hefner, Jr., et al in copending application Serial No. 790,991 filed October 24, 1985 (Attorney's Docket No. C-34,378). These resins are used to prepare thermosettable powder coatings which when cured (thermoset) provide low gloss, high impact strength coatings. Although excellent cured coatings are provided, the relatively high viscosity per unit epoxide equivalent weight of the modified advanced epoxy resins leads to difficult processability, reduced filler acceptability as well as marginal rheology on curing.

The present invention provides a specific process which results in low gloss, high impact strength epoxy resins with substantially reduced viscosity per unit epoxide equivalent weight and

resultant improvements in processability, filler acceptability and rheology.

5 One aspect of the present invention pertains to advanced epoxy resin compositions which comprise the product resulting from

10 (I) polymerizing in the presence of a catalytic quantity of at least one suitable polymerization catalyst

(A) the reaction product of

(1) a mixture comprising

15 (a) from 80 to 99, preferably from 90 to 95 percent by weight of at least one diglycidyl ether of a dihydric phenol having an epoxide equivalent weight of from 111 to 350 with the
20 epoxide equivalent weight being calculated on the basis that there are no substituent groups attached to the aromatic ring other than
25 hydrogen and the ether oxygen atoms even though the aromatic rings may in fact be substituted by groups other than hydrogen;

30 (b) from 1 to 20, preferably from 5 to 10, percent by weight of at least one diglycidyl ether of a mono-, di- or polyetherdiol; with

35 (2) at least one compound containing both a group reactive with an epoxide

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5 group and a polymerizable
ethylenically unsaturated group in an
amount of from 0.001 to 0.05,
preferably from 0.005 to 0.025,
equivalent per epoxide equivalent
contained in component (A-1); with

(B) a monomer mixture comprising

- 10 (1) at least one vinyl aromatic monomer
in an amount of from 31 to 60,
preferably from 35 to 45, percent by
weight of the combined weight of
components (A), (B) and (C);
- 15 (2) at least one compound containing both
a group reactive with an epoxide
group and a polymerizable
ethylenically unsaturated group in an
20 amount of from 0.001 to 0.05,
preferably from 0.005 to 0.025,
equivalent per epoxide equivalent
contained in component (A-1); and
optionally
- 25 (3) one or more hydroxyalkyl acrylate,
hydroxyalkyl methacrylate, alkyl
acrylate or alkyl methacrylate in an
amount of from zero to 15, preferably
30 from zero to 5, percent by weight of
the combined weight of components (B-
1) and (B-3); and

35 (II) advancing in the presence of a catalytic
quantity of at least one suitable

advancement catalyst, the product
resulting from (I) with

- 5 (C) at least one dihydric phenol in an
amount of from 0.125 to 0.8, preferably
from 0.375 to 0.5, hydroxyl equivalent
per epoxide equivalent in component (A-
1).

10 Another aspect of the present invention
pertains to advanced epoxy resin compositions which
comprise the product resulting from

- 15 (I) polymerizing in the presence of a
catalytic quantity of at least one
suitable polymerization catalyst

(A) the reaction product of

- 20 (1) at least one diglycidyl ether of a
dihydric phenol having an epoxide
equivalent weight of from 111 to 350
with the epoxide equivalent weight
being calculated on the basis that
25 there are no substituent groups
attached to the aromatic ring other
than hydrogen and the ether oxygen
atoms even though the aromatic rings
may in fact be substituted by groups
30 other than hydrogen;

- (2) at least one compound containing both
a group reactive with an epoxide
group and a polymerizable
35 ethylenically unsaturated group in an
amount of from 0.001 to 0.05,

preferably from 0.005 to 0.025,
equivalent per epoxide equivalent
contained in component (A-1); with

5 (B) a monomer mixture comprising

- 10 (1) at least one vinyl aromatic monomer
in an amount of from 31 to 60,
preferably from 35 to 45, percent by
weight of the combined weight of
components (A), (B) and (C);
- 15 (2) at least one compound containing both
a group reactive with an epoxide
group and a polymerizable
ethylenically unsaturated group in an
amount of from 0.001 to about 0.05,
preferably from 0.005 to 0.025,
equivalent per epoxide equivalent
20 contained in component (A-1); and
optionally
- 25 (3) one or more hydroxyalkyl acrylate,
hydroxyalkyl methacrylate, alkyl
acrylate or alkyl methacrylate in an
amount of from zero to 15, preferably
from zero to 5, percent by weight of
the combined weight of components (B-
1) and (B-3); and

30

35

(II) advancing in the presence of a catalytic
quantity of at least one suitable
advancement catalyst, the product
resulting from (I) with a mixture
comprising

- 5 (C) at least one dihydric phenol in an amount of from 0.125 to 0.8, preferably from 0.375 to 0.5, hydroxyl equivalent per epoxide equivalent in component (A-1) and
- 10 (D) at least one diglycidyl ether of a mono-, di- or polyetherdiol or at least one phenolic capped adduct of a diglycidyl ether of a mono-, di- or polyetherdiol or combination thereof wherein the amount of the diglycidyl ether or the amount of the diglycidyl ether contained in the capped diglycidyl ether is from 1 to 20, preferably from 5 to 10, percent by weight of components (I-A-1) and (II-D).

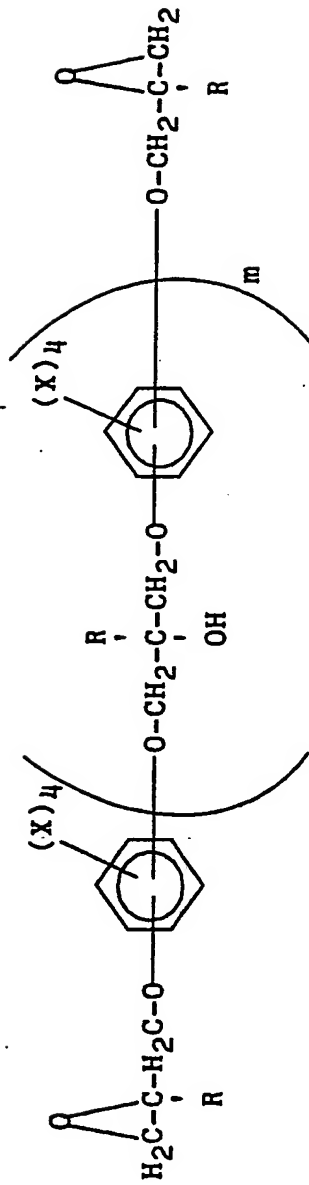
20 A further aspect of the present invention concerns thermosettable (curable) compositions prepared by combining one or more of the advanced epoxy resin compositions with a curing quantity of at least one suitable curing agent for the aforementioned advanced epoxy resin compositions.

25 Still another aspect of the present invention pertains to the products and compositions resulting from curing the aforementioned thermosettable compositions.

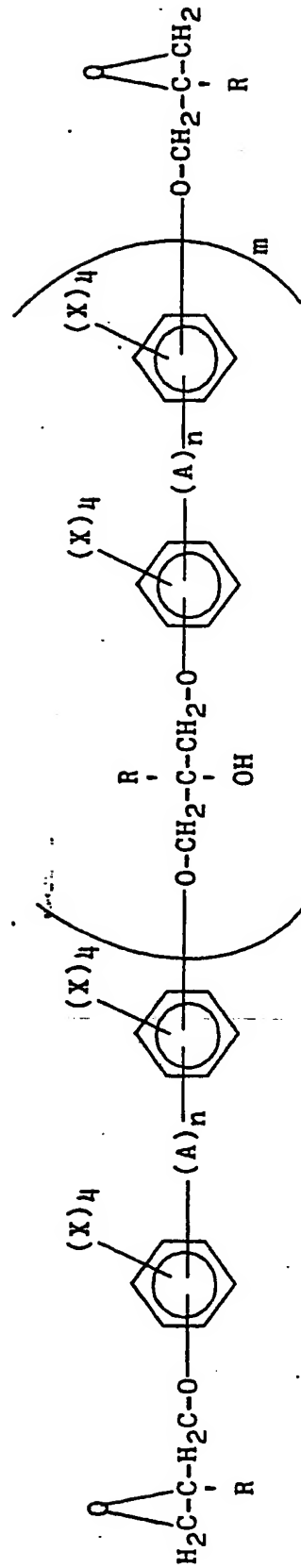
30 Suitable diglycidyl ethers of a dihydric phenol which can be employed herein include, for example, those represented by the formulas

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(I)



(II)



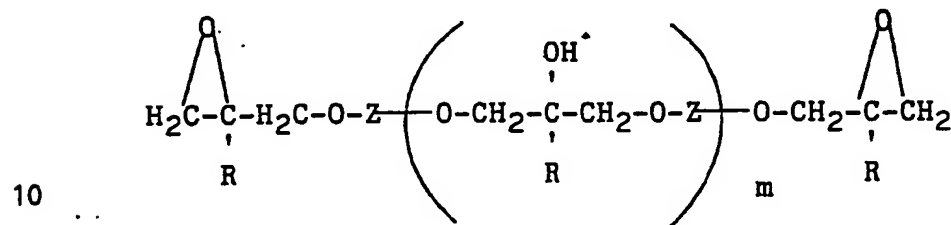
5 $\begin{array}{ccccccc} & & \text{O} & & \text{O} & & \text{O} \\ & & \parallel & & \parallel & & \parallel \\ -\text{S}-\text{S}-, & -\text{S}-, & -\text{S}-, & -\text{C}-; & \text{each X is independently hydrogen,} \\ & & \parallel & & & & \\ & & \text{O} & & & & \end{array}$

The term hydrocarbyl as employed herein means any aliphatic, cycloaliphatic, aromatic, aryl substituted aliphatic, or aliphatic substituted aromatic groups. Likewise, the term hydrocarbyloxy group means a hydrocarbyl group having an oxygen linkage between it and the object to which it is attached.

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Suitable diglycidyl ethers of a mono-, di- or polyetherdiol which can be employed herein include, for example, those represented by the formula

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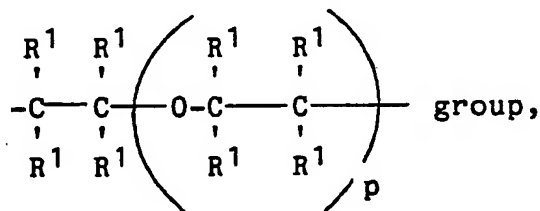
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(III)

wherein Z is a hydrocarbyl group containing from 1 to 12 carbon atoms or a

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wherein each R^1 is independently hydrogen or a hydrocarbyl group having from 1 to 6 carbon atoms, and p has a value from 1 to 100, preferably from 1 to 25; m and R are as hereinbefore defined. The term mono-, di- or polyetherdiol is intended to also include aliphatic or cycloaliphatic diols or mixtures of said diols.

30

Suitable diglycidyl ethers of a mono-, di- or polyetherdiol which can be employed herein include, for example, the diglycidyl ethers of ethylene glycol, propylene glycol, diethylene glycol, dipropylene glycol, triethylene glycol, tripropylene glycol,

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butylene glycol, polyethylene glycols, polypropylene glycols, polybutylene glycols, 1,12-dihydroxydodecane, 1,2-dihydroxyundecane, 1,6-hexanediol, 1,3-dihydroxy-2,2-dimethyl propane, and mixtures thereof. Most
5 preferred are the diglycidyl ethers of ethylene glycol, propylene glycol, polyethylene glycol and polypropylene glycol.

The diglycidyl ethers of a mono-, di- or
10 polyetherdiol may be converted to a phenolic capped adduct suitable for use herein via reaction with a stoichiometric excess of a dihydric phenol such as those represented by formulas VI and VII. An
15 equivalent ratio of 1:2 or more epoxide groups to phenolic hydroxyl groups is used to prepare the phenolic capped adduct. An equivalent ratio of 1:2.5 to 1:5 is most preferred. The phenolic capped adduct is preferably prepared using an advancement catalyst
20 with reaction times and temperatures delineated herein for advancement reactions.

Suitable compounds which contain both a group reactive with an epoxide group and a polymerizable
25 ethylenically unsaturated group include compounds wherein said group reactive with an epoxide group is a compound containing a carboxylic acid, hydroxyl or amido group. Suitable compounds which contain both a group reactive with an epoxide group and a
30 polymerizable ethylenically unsaturated group include, for example, the acrylic acids, such as acrylic acid and methacrylic acid; the monoesters of α,β -unsaturated dicarboxylic acids, such as monomethyl maleate and monobutylfumarate; the alkenylphenols such as p-
35 isopropenylphenol, propenylphenol and m-vinylphenol; the hydroxyalkyl acrylates such as 2-hydroxyethyl

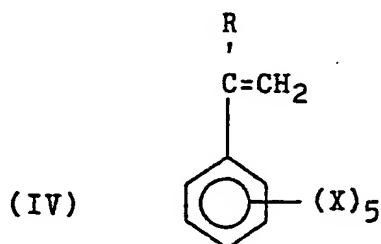
acrylate, 2-hydroxyethyl methacrylate, 2-hydroxypropyl acrylate, 2-hydroxypropyl methacrylate and the acrylamides such as methacrylamide and acrylamide, any combination thereof and the like. Most preferred as
5 the compound containing a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group is methacrylic acid.

The prereaction (step A) of the diglycidyl
10 ether of a dihydric phenol or the diglycidyl ether of a dihydric phenol combined with the diglycidyl ether of a diol and a compound which contains both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group is performed at a
15 temperature of from 75 to 200°C, preferably from 140 to 160°C for from 15 minutes to 150 minutes, preferably for from 30 minutes to 60 minutes. The prereaction step times and temperatures vary as a function of the type
20 of compound which contains both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group that is used.

A catalyst may optionally be employed to
25 facilitate reaction of the group reactive with an epoxide group and the epoxide group. Generally, a catalyst is not required and, furthermore, is not desired when said group reactive with an epoxide group is -COOH. It may, however, be beneficial to use a
30 catalyst when said group reactive with an epoxide group is, for example, -OH. Typical of such catalysts useful for this purpose are the advancement catalysts described herein.

35 Suitable vinyl aromatic monomers which can be employed as component (B-1) in the copolymerization

reaction step with the prereaction product of a diglycidyl ether of a dihydric phenol or the diglycidyl ether of a dihydric phenol combined with the diglycidyl ether of a diol and a compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group include those represented by the formula



wherein R and X are as hereinbefore defined.

Representative of the vinyl aromatic monomers which can be employed herein are styrene, chlorostyrenes, methylstyrenes, t-butylstyrenes, α -methylstyrene, methoxystyrenes, and mixtures thereof. Most preferred as the vinyl aromatic monomer is styrene.

Suitable hydroxyalkyl acrylates or methacrylates, alkyl acrylates or methacrylates or mixtures thereof which can be employed as component (B-3) in the copolymerization reaction step with the prereaction product of a diglycidyl ether of a dihydric phenol or the diglycidyl ether of a dihydric phenol combined with the diglycidyl ether of a diol and a compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group may be chosen according to structure and amount so as to

effect the final properties of a cured composition such as, for example, a powder coating. Small amounts (0.25 to 2 percent by weight based on total weight of monomer mixture used) of a hydroxyalkyl acrylate or meth-
5 acrylate are used to increase adhesion of the powder coating to metal substrates. Larger amounts (2.1 to 15 percent by weight based on total weight of monomer mixture used) of a hydroxyalkyl acrylate or
10 methacrylate increase the gloss of the powder coating. The alkyl acrylates or methacrylates, especially those possessing 8 or more carbon atoms, are used in small amounts (1 to 5 percent by weight based on total weight of monomer mixture used) to decrease the gloss of the
15 powder coating. Larger amounts (5.1 to 15 percent by weight based on total weight of monomer mixture used) of certain alkyl acrylates or methacrylates can be used to also impart modified texture to the powder coating. Combinations of said acrylates and methacrylates may
20 also be used. Specific hydroxyalkyl acrylates or methacrylates, alkyl acrylates or methacrylates which can optionally be employed herein include those represented by the formula



wherein R is as hereinbefore defined and Q is a
30 monovalent hydrocarbyl group having from one to 25 carbon atoms or a hydroxyalkyl group having from two to 25 carbon atoms and may be branched, cyclic or polycyclic. Representative of the hydroxyalkyl acrylates or methacrylates, alkyl acrylates or
35 methacrylates or mixtures thereof which can optionally be employed herein are 2-hydroxyethyl acrylate, 2-

hydroxyethyl methacrylate, 2-hydroxypropyl acrylate, cyclohexyl acrylate, lauryl methacrylate, stearyl acrylate, mixtures thereof and the like.

5 Suitable free radical forming catalysts which
can be employed in the copolymerization reaction step
with the prereaction product of a diglycidyl ether of a
dihydric phenol or the diglycidyl ether of a dihydric
phenol combined with the diglycidyl ether of a diol and
10 a compound containing both a group reactive with an
epoxide group and a polymerizable ethylenically unsaturated group include the azo and diazo compounds as well
as the organic peroxides and hydroperoxides. Suitable
free radical forming catalysts include, for example,
15 2,2'-azobisisobutyronitrile, 2,2'-azobis(2,4-dimethyl-
4-methoxyvaleronitrile), 1-t-butylazo-1-cyanocyclo-
hexane, t-butylperbenzoate, t-butyl-peroctoate,
t-butylhydroperoxide, di-t-butylperoxide, dicumyl-
peroxide, cumene hydroperoxide, and mixtures thereof.
20 An amount of from 1.0 to 5.0, preferably from 2.0 to
3.0 percent by weight, based on total weight of monomer
mixture used, of at least one free radical forming
catalyst is employed.

25 The copolymerization reaction of the
prereaction product (A) of a diglycidyl ether of a
dihydric phenol or the diglycidyl ether of a dihydric
phenol combined with the diglycidyl ether of a diol and
30 a compound containing both a group reactive with an
epoxide group and a polymerizable ethylenically unsaturated group with a monomer mixture consisting of (B-1)
a vinyl aromatic monomer, (B-2) a compound containing
both a group reactive with an epoxide group and a
35 polymerizable ethylenically unsaturated group, and
optionally, (B-3) a hydroxyalkyl acrylate or

methacrylate, an alkyl acrylate or methacrylate, or a mixture thereof may be completed using a variety of reaction sequences. Generally, the monomer feed (B) containing a free radical forming polymerization catalyst is added to the prereaction product (A) over a period of from 45 minutes to 150 minutes, preferably from 75 minutes to 120 minutes while maintaining a reaction temperature of from 125° to 175°C, preferably from 140° to 160°C. A post reaction of from 30 minutes to 150 minutes, preferably from 45 minutes to 90 minutes is completed after completion of the monomer feed addition.

It is necessary to maintain an inert atmosphere throughout the copolymerization reaction. This is achieved by blanketing the reaction mixture with nitrogen, argon or some other inert gas. Adequate stirring is required to intimately mix and disperse the reactants.

In an equally preferred process of the present invention, the free radical forming catalyst may be removed as a component of the monomer feed and added to the reaction mixture separately. If this is done, it is generally desirable to maintain concurrent addition of the free radical forming catalyst and the remaining monomer mixture (B-1, B-2 and, optionally, B-3). The rate of this concurrent addition should be adjusted such that an excess of unpolymerized monomer mixture does not accumulate.

As a further embodiment of the present invention, a portion of free radical forming catalyst may be added to the reaction mixture at the end of the monomer mixture addition, more preferably 15 minutes to

120 minutes after completion of the monomer mixture addition. This is done if unpolymerized monomer mixture is present and allows for completion of the copolymerization reaction.

5

When a diglycidyl ether of a dihydric phenol combined with the diglycidyl ether of a diol is used to prepare the prereaction product (A) the diglycidylether of a diol (A-1-b) may be added to the reaction mixture at any time but preferably is added simultaneous with the diglycidyl ether of a dihydric phenol.

10

The advancement reaction of the aforementioned copolymerization product of the prereaction product (step A) of the diglycidyl ether of a dihydric phenol combined with the diglycidyl ether of a diol and a compound which contains both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group and the monomer mixture (step B) with a dihydric phenol is performed in the presence of an advancement catalyst. Suitable dihydric phenols include, for example, those represented by the formulas

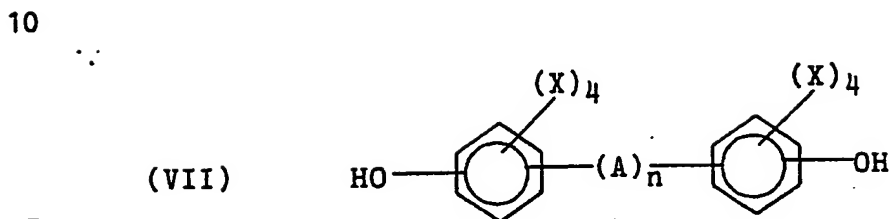
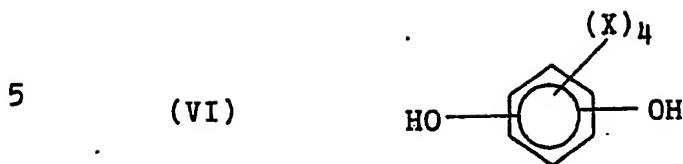
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wherein X, A and n are as hereinbefore defined. Representative of the bisphenols are resorcinol, hydroquinone, catechol, bisphenol A (4,4'-isopropylidenediphenol), bis(4,4'-dihydroxyphenyl)methane, 2,2'-bis(4-hydroxyphenyl)pentane, 3,3',5,5'-tetramethyl-4,4'-dihydroxydiphenyl, 4,4'-dihydroxydiphenyl sulfone, 4,4'-dihydroxybenzophenone, 3,3',5,5'-tetrabromo-4,4'-isopropylidenediphenol, 4,4'-bis(p-hydroxyphenyl)diphenyl ether, 4,4'-dihydroxydiphenyl sulfide, and mixtures thereof. Most preferred as the bisphenol is bisphenol A.

30 Suitable advancement catalysts which can be employed in the process of the present invention include most any catalyst which will catalyze the reaction between a vicinal epoxy group and a phenolic hydroxyl group. Such catalysts include, for example, those disclosed in U.S. Patent Nos. 3,306,872; 3,341,580; 3,379,684; 3,477,990; 3,547,881; 3,637,590;

3,843,605; 3,948,855; 3,956,237; 4,048,141; 4,093,650;
4,131,633; 4,132,706; 4,171,420; 4,177,216.

Particularly suitable catalysts are the
5 quaternary phosphonium and ammonium compounds such as,
for example, ethyltriphenylphosphonium chloride,
ethyltriphenylphosphonium bromide, ethyltriphenyl-
phosphonium iodide, ethyltriphenylphosphonium acetate,
ethyltriphenylphosphonium acetate-acetic acid complex,
10 ethyltriphenylphosphonium phosphate, tetrabutyl-
phosphonium chloride, tetrabutylphosphonium bromide,
tetrabutylphosphonium iodide, tetrabutylphosphonium
acetate, tetrabutylphosphonium acetate-acetic acid
complex, butyltriphenylphosphonium tetrabromo-
15 bisphenate, butyltriphenylphosphonium bisphenate,
butyltriphenylphosphonium bicarbonate, benzyltri-
methylammonium chloride, tetramethylammonium hydroxide,
and mixtures thereof.

20 The advancement reaction of the aforemen-
tioned copolymerization product of the prereaction
product (step A) of the diglycidyl ether of a dihydric
phenol and a compound which contains both a group
25 reactive with an epoxide group and a polymerizable
ethylenically unsaturated group and the monomer mixture
(step B) with a dihydric phenol (formulas VI, VII) and
a diglycidyl ether of a diol or a phenolic capped
adduct of a diglycidyl ether of a diol is performed in
30 the presence of an advancement catalyst using methods
hereinbefore delineated.

It is also operable, although less preferred,
to perform the advancement reaction of the aforemen-
35 tioned copolymerization product of the prereaction
product of the diglycidyl ether of a dihydric phenol

combined with the diglycidyl ether of a diol and a compound which contains both a group reactive with an epoxide group (Step A) and a polymerizable ethylenically unsaturated group and the monomer mixture (Step
5 B) with a dihydric phenol (formulas VI or VII) and a diglycidyl ether of a diol or a phenolic capped adduct of a diglycidyl ether of a diol in the presence of an advancement catalyst using methods hereinbefore delineated. In this particular process configuration, the
10 previously specified prerequisite amount of diglycidyl ether of a diol is thus distributed between the pre-reaction and the advancement reaction steps in any proportion desired.

15 After completion of either the copolymerization reaction (I) or the advancement reaction (II), it is generally beneficial, although not required, to subject the reaction product to a vacuum stripping
20 step. This is accomplished by pulling a vacuum on the reactor, thus removing and condensing any materials which volatilize from the reaction product.

25 In a variation on this vacuum stripping step, various modifying agents or additives, such as, for example, flow control agents, gloss control agents, pigments, texture control additives, air release agents, mixtures thereof and the like may be added to the reaction product prior to the vacuum stripping
30 step. This allows for removal of any volatile components contributed by said additives.

35 The powder coating formulations of the present invention are prepared using the aforesaid epoxy resin compositions and a suitable curing agent therefor. Said curing agents should be substantially non-

sintering and exhibit latency at the formulating temperatures and conditions employed. Suitable such curing agents are described in the Handbook of Epoxy Resins, by Lee and Neville, 1967, McGraw-Hill.

5 Representative of these curing agents are the boron trifluorideamine complexes, polycarboxylic acids or anhydrides, the guanamines, guanidines, hydrazines, dihydrazides, melamines, substituted ureas and
10 biguanides, such as, dicyandiamide or dicyandiamide derivatives. Most preferred as the curing agent is dicyandiamide.

15 Epoxy polyester hybrid powder coating formulations of the present invention are prepared using the aforesaid epoxy resin compositions and a suitable carboxylic acid and/or hydroxyl functional polyester as the curing agent.

20 Formulating methods well known to the prior art are employed to prepare the powder coating formulations of the present invention. Preparation of typical epoxy resin based powder coating formulations are described in Fundamentals of Powder Coating by Miller
25 and Taft, 1974, Society of Manufacturing Engineers, Dearborn, Michigan.

30 In the general method of preparation, the solid epoxy resin product is flaked or ground then dry mixed or blended with a non-sintering curing agent and optionally, one or more curing agent accelerators or catalysts, particulate fillers, pigments, flow control agents, gloss control additives, texture control additives and air release agents. The dry mixed product is
35 then hot melt blended typically by use of a kneading-type extruder. The extruded product passes through

chilled rollers and is then recovered and crushed to a rough powder. Further grinding to a fine powder is accomplished via use of a high speed hammer mill or other type of grinding equipment. The resulting fine powder is subjected to a size classification step to recover the desired range of product particle size. The desired product size distribution for the product may vary depending on the intended end use of the product, but generally, sizes between 80 mesh to 325 mesh are most desired. Well known methods that are suitable for use in size classifying powder coating formulations include screening and air classification.

The resulting powder coating formulation is applied to the substrate to be coated using methods well known to the prior art. These methods are delineated in detail by the aforementioned Miller and Taft reference and include powder dusting, fluidized bed processes, electrostatic powder spraying, electrostatic fluidized bed processes, and others.

The powder coated article is cured using methods and conditions well known to the prior art. This typically involves heating in an oven for an amount of time sufficient to complete the cure. When dicyandiamide is the curing agent and 2-methylimidazole is the curing agent accelerator used with the epoxy resin compositions of the present invention, curing times of 5 minutes to 30 minutes at a reaction temperature of from 150 to 220°C are generally sufficient.

The powder coating formulation optionally, although preferably, contains one or more curing agent accelerators or catalysts. Suitable such curing agent

accelerators or catalysts are described in the aforementioned Handbook of Epoxy Resins and Fundamentals of Powder Coating references. Representative of these curing agent accelerators or catalysts are the amino substituted pyridines, imidazoles, metallic salts, tertiary amines, phenols, and mixtures thereof. Most preferred as the curing agent accelerator for use with a dicyandiamide curing agent is 2-methylimidazole.

10 The powder coating formulation optionally, although preferably, contains one or more particulate fillers. Fillers are used in powder coatings for a wide range of purposes, primary of which is economic, i.e. as a less expensive diluent. Other properties
15 imparted by fillers can include one or more of the following: handling and processing properties, impact modification, dimensional stability, moisture and chemical resistance, flame resistance, modified thermal conductivity, modified electrical properties, modified
20 rheology, color modification and texture modification. Suitable such fillers are described in Non-Fibrous Fillers for Epoxy Resin Formulations presented at the 7th Electrical Insulation Conference, Chicago, Illinois, October 15-19, 1967 by D. A. Shimp.
25 Representative of these fillers are barytes (BaSO_4), titanium dioxide, carbon black, silica flour, calcium carbonate, and mixtures thereof. The particle size distribution, shape, chemical composition, surface area
30 and use level, i.e. resin to filler ratio, can be adjusted singularly or collectively to change the resultant cured powder coating. Simple preliminary experiments within the normal capability of those
35 skilled in the art are ordinarily performed to aid in filler choice.

The powder coating formulation optionally contains one or more pigments. Said pigments are typically used to add color to the cured powder coating. Suitable such pigments are described in
5 Pigments for Colouring Epoxy Powder Coatings by Maltman and Deverell-Smith in Pigment and Resin Technology, November 1973, pp. 15-19.

The powder coating formulation optionally,
10 although preferably, contains one or more flow control agents. Flow control agents are used in powder coatings to adjust the rheological properties of the total powder coating formulation thus insuring uniform coating film thickness, wet-out and coating of edges.
15 Suitable such flow control agents are described in Acrylic Flow Control Agents for the Coating Industry by Skora in Polymers Paint and Colour Journal, September 5, 1979, pp. 867-870. Most preferred as the flow
20 control agents are the polyacrylates such as, for example, ethyl acrylate and 2-ethylhexyl acrylate copolymer, and poly(butyl acrylate).

The powder coating formulation optionally
25 contains one or more texture control additives. Texture control additives are used in powder coatings to modify the surface characteristics of the cured powder coating. Materials which provide smooth or rough surface finishes may be employed. Glass
30 microspheres, metal powders and polymeric powders are examples of the types of additives capable of modifying the powder coating surface to a textured finish.

The powder coating formulation optionally
35 contains one or more air release agents. Said agents are used in powder coatings to alleviate surface

defects, such as pinholes in the cured powder coating, induced by air entrainment. A most preferred air release agent is benzoin, as described in Surface Coatings, Vol. 2 - Paints and Their Application by The Oil and Colour Chemists' Association, Australia, published by Chapman and Hall, 1984, p. 598.

The powder coating formulation optionally contains one or more gloss control additives. Gloss control additives are used to reduce the high degree of reflected light from the typical cured epoxy resin surface. Suitable such gloss control agents are certain amorphous silicas, silicic acid and the curing agent system consisting of a salt of a polycarboxylic acid and a cyclic amidine as taught by U.S. Patent No. 3,947,384.

Other additives or adjuvants may be incorporated into the powder coating formulations of the present invention for their known and intended use therein. One such additive is a slip aid, as described in the aforementioned Surface Coatings reference.

The cured product of the present invention is a powder coating over a substrate such as steel which provides a smooth finish, low gloss surface with high impact strength.

The following examples are illustrative of the present invention and are not to be construed as to limiting the scope thereof in any manner.

EXAMPLE 1

Polystyrene Modified Epoxy Resin with 5% of the Diglycidyl Ether of Bisphenol A Substituted with

Polypropylene Glycol Diglycidyl Ether During the
Prereaction

5 A commercial grade (D.E.R.® 383, The Dow
Chemical Company) diglycidyl ether of bisphenol A (575
g, 3.144 epoxide equivalents) having an epoxide equiv-
alent weight (EEW) of 182.9, a commercial grade
(D.E.R.® 732, The Dow Chemical Company) polypropylene
glycol diglycidyl ether (49.14 g, 0.166 epoxide
10 equivalent) having an EEW of 296 and glacial
methacrylic acid (2.86 g, 0.033 mole) were charged to a
2-liter, 5-neck round bottom flask fitted with a water-
cooled condenser, nitrogen purge, addition funnel,
thermometer and stirrer. Heating to 150°C was initiated
15 on the stirred solution. Forty-four minutes later the
reactor temperature reached 145°C and a nitrogen purge
was initiated. Two minutes later the 150°C temperature
was reached. The first aliquot of monomer mixture, a
mixture of styrene (249.65 g, 2.39 moles), glacial
20 methacrylic acid (2.15 g, 0.025 mole) and tertiary
butyl perbenzoate (6.24 g, 0.031 mole) was charged to
the addition funnel and added dropwise to the stirred
solution. After forty-four minutes, the addition of
25 the first aliquot of monomer mixture was complete and
the second aliquot, a mixture of styrene (249.65 g,
2.39 moles), glacial methacrylic acid (2.15 g, 0.025
mole) and tertiary butyl perbenzoate (6.24 g, 0.031
mole), was charged to the addition funnel and added
30 dropwise to the stirred solution. The reactor
temperature was maintained between 145°C and 155°C
throughout the monomer mixture addition. Forty-seven
minutes later the addition of the second aliquot of
monomer mixture was complete. Post-reaction of the
35 stirred solution was continued for thirty minutes at

150°C. After thirty minutes at 150°C, tertiary butyl perbenzoate (1.25 g, 0.0062 mole) was charged to the stirred solution. After an additional thirty minutes at 150°C, a vacuum was drawn to remove unreacted styrene and other lights. During the vacuum stripping, 2.0 ml of unreacted styrene and other lights were recovered in an overhead cold trap. After thirty minutes of vacuum stripping at 150°C, bisphenol A (162.8 g, 0.71 mole) was charged to the stirred solution. Three minutes later, the reactor temperature had dropped to 133°C with complete dissolution of the bisphenol A in the resin solution. An advancement catalyst (ethyltriphenylphosphonium acetate-acetic acid complex, 1 g, 70% in methanol) was added to the reactor which was then reheated to 150°C. Seven minutes later, the reactor reached 150°C and heating was discontinued. A maximum exotherm of 180°C resulted six minutes later. The reactor was allowed to cool to 175°C where it was held for one hour after the maximum exotherm. The polystyrene modified epoxy resin product was recovered as a pale-yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 721.

25 EXAMPLE 2

30 Polystyrene Modified Epoxy Resin with 5% of the Diglycidyl Ether of Bisphenol A Substituted with Polypropylene Glycol Diglycidyl Ether After Monomer Mixture Addition

A commercial grade (D.E.R.* 383, The Dow Chemical Company) diglycidyl ether of bisphenol A (575 g, 3.144 epoxide equivalents) having an epoxide equivalent weight (EEW) of 182.9 and glacial methacrylic acid (2.86 g, 0.033 mole) were charged to a

2-liter, 5-neck round bottom flask fitted with a water-cooled condenser, nitrogen purge, addition funnel, thermometer and stirrer. Heating to 150°C was initiated on the stirring solution. Forty-seven minutes later, the reactor temperature reached 145°C and a nitrogen purge was initiated. Three minutes later, the 150°C temperature was reached. The first aliquot of monomer mixture, a mixture of styrene (249.65 g, 2.39 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6.24 g, 0.031 mole) was charged to the addition funnel and added dropwise to the stirred solution. After fifty minutes, the addition of the first aliquot of monomer mixture was complete and the second aliquot, a mixture of styrene (249.65 g, 2.39 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6.24 g, 0.031 mole), was charged to the addition funnel and added dropwise to the stirred solution. The reactor temperature was maintained between 145°C and 155°C throughout the monomer mixture addition. Forty-three minutes later, the addition of the second aliquot of monomer mixture was complete. Post-reaction of the stirred solution was continued for thirty minutes at 150°C. After fifteen minutes at 150°C, tertiary butyl perbenzoate (1.25 g, 0.0062 mole) was charged to the stirred solution. After an additional fifteen minutes at 150°C a vacuum was drawn to remove unreacted styrene and other lights. During the vacuum stripping, 7.5 ml of unreacted styrene and other lights were recovered in an overhead cold trap. After sixty minutes of vacuum stripping at 150°C, a commercial grade (D.E.R.* 732, The Dow Chemical Company) polypropylene glycol diglycidyl ether (49.14 g, 0.166 epoxide equivalent) having an EEW of 296 and bisphenol A (162.8 g, 0.71 mole) were

charged to the stirred solution. Six minutes later, the reactor temperature had dropped to 130°C with complete dissolution of the bisphenol A in the resin solution. An advancement catalyst
5 (ethyltriphenylphosphonium acetate-acetic acid complex, 1 g, 70% in methanol) was added to the reactor which was then reheated to 150°C. Eight minutes later, the reactor reached 150°C and heating was discontinued. A maximum exotherm of 180°C resulted seven minutes
10 (420 s) later. The reactor was allowed to cool to 175°C where it was held for one hour after the maximum exotherm. The polystyrene modified epoxy resin product was recovered at room temperature (25°C) as a pale-
15 yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 737.

EXAMPLE 3

Polystyrene Modified Epoxy Resin with 5% of the Diglycidyl Ether of Bisphenol A Substituted with Polypropylene Glycol Diglycidyl Ether During the Advancement Step Post Reaction

A commercial grade (D.E.R.[®] 383, The Dow
25 Chemical Company) diglycidyl ether of bisphenol A (575 g, 3.144 epoxide equivalents) having an epoxide equivalent weight (EEW) of 182.9 and glacial methacrylic acid (2.86 g, 0.033 mole) were charged to a
30 2-liter, 5-neck round bottom flask fitted with a water-cooled condenser, nitrogen purge, addition funnel, thermometer and stirrer. Heating to 150°C was initiated on the stirring solution. Forty-two minutes later, the reactor temperature reached 145°C and a nitrogen purge
35 was initiated. Three minutes later, the 150°C temperature was reached. The first aliquot of monomer

mixture, a mixture of styrene (249.65 g, 2.39 moles),
glacial methacrylic acid (2.15 g, 0.025 mole) and
tertiary butyl perbenzoate (6.24 g, 0.031 mole) was
charged to the addition funnel and added dropwise to
5 the stirred solution. After forty-five minutes, the
addition of the first aliquot of monomer mixture was
complete and the second aliquot, a mixture of styrene
(249.65 g, 2.39 moles), glacial methacrylic acid (2.15
10 g, 0.025 mole) and tertiary butyl perbenzoate (6.24 g,
0.031 mole), was charged to the addition funnel and
added dropwise to the stirred solution. The reactor
temperature was maintained between 145°C and 155°C
throughout the monomer mixture addition. Forty-three
15 minutes later, the addition of the second aliquot of
monomer mixture was complete. Post-reaction of the
stirred solution was continued for fifteen minutes at
150°C. After fifteen minutes at 150°C, tertiary butyl
perbenzoate (1.25 g, 0.0062 mole) was charged to the
20 stirred solution. After an additional forty-five
minutes at 150°C a vacuum was drawn to remove unreacted
styrene and other lights. During the vacuum stripping,
2.7 ml of unreacted styrene and other lights were
recovered in an overhead cold trap. After forty-two
25 minutes of vacuum stripping at 150°C, bisphenol A (162.8
g, 0.71 mole) was charged to the stirred solution.
Three minutes later, the reactor temperature had
dropped to 130°C with complete dissolution of the
30 bisphenol A in the resin solution. An advancement
catalyst (ethyltriphenylphosphonium acetate·acetic
acid complex, 1 g, 70% in methanol) was added to the
reactor which was then reheated to 150°C. Nine minutes
later, the reactor reached 150°C and heating was
35 discontinued. A maximum exotherm of 181°C resulted
seven minutes later. The reactor was allowed to cool

to 175°C where it was held for forty-five minutes after the maximum exotherm. A commercial grade (D.E.R.® 732, The Dow Chemical Company) of a polypropylene glycol diglycidyl ether of (49.14 g, 0.166 epoxide equivalent) was added to the advanced resin mixture. The reactor was held at 175°C for fifteen minutes following the polypropylene glycol diglycidyl ether addition. The polystyrene modified epoxy resin product was recovered at room temperature (25°C) as a pale-yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 733.

EXAMPLE 4

15 Polystyrene Modified Epoxy Resin with 5% of the Diglycidyl Ether of Bisphenol A Substituted with Bisphenol A Capped Polypropylene Glycol Diglycidyl Ether After Monomer Mixture Addition

20 A. Synthesis of Phenolic Capped Aliphatic Epoxy Resin

A portion of a commercial grade (D.E.R.® 732, The Dow Chemical Company) diglycidyl ether of polypropylene glycol (59.20 g, 0.20 epoxide equivalent) having an epoxide equivalent weight (EEW) of 296 was added to a reactor and heated to 150°C with stirring under a nitrogen atmosphere. Bisphenol A (68.49 g, 0.60 hydroxyl equivalent) was added to the reactor followed by cooling to 100°C. At this time, ethyltri-phenylphosphonium bromide-acetic acid complex catalyst (0.192 g, 0.15 percent by weight) was added to the reactor followed by heating to 180°C over a 16 minute period. Once the 180°C temperature was achieved, the reactor was allowed to cool over a 10 minute period to 150°C then held at this temperature for 64 minutes. The

product was recovered as a light yellow colored, transparent solid at room temperature (25°C) with a residual percent epoxide content of 0.34 (12,647 EEW).

B. Synthesis of a Polystyrene Modified Epoxy Resin
5 Utilizing a Portion of the Phenolic Capped
Aliphatic Epoxy Resin

A commercial grade (D.E.R.* 383, The Dow Chemical Company) diglycidyl ether of bisphenol A
10 (489.72 g, 2.679 epoxide equivalents) having an epoxide equivalent weight (EEW) of 182.8 and glacial methacrylic acid (2.86 g, 0.033 mole) were charged to a 2-liter, 5-neck round bottom flask fitted with a water-cooled condenser, nitrogen purge, addition funnel,
15 thermometer and stirrer. Heating to 150°C was initiated on the stirred solution. Thirty-two minutes later, the reactor temperature reached 145°C and a nitrogen purge was initiated. Two minutes later, the 150°C temperature
20 was reached. The first aliquot of monomer mixture, a mixture of styrene (212.58 g, 2.04 moles), glacial methacrylic acid (1.82 g, 0.021 mole) and tertiary butyl perbenzoate (5.315 g, 0.027 mole) was charged to the addition funnel and added dropwise to the stirred
25 solution. After forty-six minutes, the first aliquot of monomer mixture addition was complete and the second aliquot, a mixture of styrene (212.58 g, 2.04 moles), glacial methacrylic acid (1.82 g, 0.021 mole) and tertiary butyl perbenzoate (5.315 g, 0.027 mole), was
30 charged to the addition funnel and added dropwise to the stirred solution. The reactor temperature was maintained between 145°C and 155°C throughout the monomer mixture addition. Forty-two minutes later, the
35 second aliquot of monomer mixture addition was complete. Post-reaction of the stirred solution was

continued for thirty minutes at 150°C. After thirty minutes at 150°C, tertiary butyl perbenzoate (1.06 g, 0.0054 mole) was charged to the stirred solution. After an additional thirty minutes at 150°C, a vacuum
5 was drawn to remove unreacted styrene and other lights. During the vacuum stripping, 4.85 ml of unreacted styrene and other lights were recovered in an overhead cold trap. After thirty minutes of vacuum stripping at
10 150°C, 90 g of the phenolic capped aliphatic epoxy resin product (part A) was charged to the stirred solution concurrently with bisphenol A (90.28 g, 0.394 mole). Ten minutes later, the reactor temperature had dropped to 134°C with complete dissolution of the bisphenol A in
15 the resin solution. An advancement catalyst (ethyltriphenylphosphonium acetate-acetic acid complex, 0.84 g, 70% in methanol) was added to the reactor which was then reheated to 150°C. Six minutes later, the reactor reached 150°C and heating was
20 discontinued. A maximum exotherm of 180°C resulted five minutes later. The reactor was allowed to cool to 175°C where it was held for one hour after the maximum exotherm. The polystyrene modified epoxy resin product was recovered at room temperature (25°C) as a pale-
25 yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 738.

EXAMPLE 5

30 Polystyrene Modified Epoxy Resin with 10% of the Diglycidyl Ether of Bisphenol A Substituted with Polypropylene Glycol Diglycidyl Ether During the Prereaction

35 A commercial grade (D.E.R.[®] 383, The Dow Chemical Company) diglycidyl ether of bisphenol A

(540.99 g, 2.979 epoxide equivalents) having an epoxide equivalent weight (EEW) of 181.6, a commercial grade (D.E.R.[®] 732, The Dow Chemical Company) polypropylene glycol diglycidyl ether (97.98 g, 0.331 epoxide equivalent) having an EEW of 296 and glacial methacrylic acid (2.86 g, 0.033 mole) were charged to a 2-liter, 5-neck round bottom flask fitted with a water-cooled condenser, nitrogen purge, addition funnel, thermometer and stirrer. Heating to 150°C was initiated on the stirred solution. Forty-seven minutes later, the reactor temperature reached 145°C and a nitrogen purge was initiated. Five minutes later, the 150°C temperature was reached. The first aliquot of monomer mixture, a mixture of styrene (255.59 g, 2.45 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6.39 g, 0.032 mole) was charged to the addition funnel and added dropwise to the stirred solution. After thirty-eight minutes, the first aliquot of monomer mixture addition was complete and the second aliquot, a mixture of styrene (255.59 g, 2.45 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6.39 g, 0.032 mole), was charged to the addition funnel and added dropwise to the stirred solution. The reactor temperature was maintained between 145°C and 155°C throughout the monomer mixture addition. Fifty-one minutes later, the second aliquot of monomer mixture addition was complete. Post-reaction of the stirred solution was continued for thirty-two minutes at 150°C. After thirty-two minutes at 150°C, tertiary butyl perbenzoate (1.28 g, 0.0064 mole) was charged to the stirred solution. After twenty-eight minutes at 150°C, a vacuum was drawn to remove unreacted styrene and other lights. During the vacuum stripping, 7.0 ml of

unreacted styrene and other lights were recovered in an overhead cold trap. After thirty minutes of vacuum stripping at 150°C, bisphenol A (158.97 g, 0.69 mole) was charged to the stirred solution. Four minutes later, the reactor temperature had dropped to 134°C with complete dissolution of the bisphenol A in the resin solution. An advancement catalyst (ethyltriphenylphosphonium acetate-acetic acid complex, 1.0 g, 70% in methanol) was added to the reactor which was then reheated to 150°C. Seven minutes later, the reactor reached 150°C and heating was discontinued. A maximum exotherm of 176°C resulted four minutes later. The reactor was allowed to cool to 175°C where it was held for one hour after the maximum exotherm. The polystyrene modified epoxy resin product was recovered at room temperature (25°C) as a pale-yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 740.

COMPARATIVE EXPERIMENT A

Polystyrene Modified Epoxy Resin - Low Gloss Standard

A commercial grade (D.E.R.[®] 383, The Dow Chemical Company) diglycidyl ether of bisphenol A (600 g, 3.31 epoxide equivalents) having an epoxide equivalent weight (EEW) of 181.6 and glacial methacrylic acid (2.86 g, 0.033 mole) were charged to a 2-liter, 5-neck round bottom flask fitted with a water-cooled condenser, nitrogen purge, addition funnel, thermometer and stirrer. Heating to 150°C was initiated on the stirred solution. Forty-two minutes later, the reactor temperature reached 145°C and a nitrogen purge was initiated. Three minutes later, the 150°C temperature was reached. The first aliquot of monomer mixture, a

mixture of styrene (240 g, 2.3 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6 g, 0.03 mole) was charged to the addition funnel and added dropwise to the stirred solution. After forty-seven minutes, the addition of the first aliquot of monomer mixture was complete and the second aliquot, a mixture of styrene (240 g, 2.3 moles), glacial methacrylic acid (2.15 g, 0.025 mole) and tertiary butyl perbenzoate (6 g, 0.03 mole), was charged to the addition funnel and added dropwise to the stirred solution. The reactor temperature was maintained between 145°C and 155°C throughout the monomer mixture addition. Forty minutes later, the addition of the second aliquot of monomer mixture was complete. Post-reaction of the stirred solution was continued for thirty minutes at 150°C. After thirty minutes at 150°C, tertiary butyl perbenzoate (1.2 g, 0.006 mole) was charged to the stirred solution. A slight exotherm to 153°C was noted. After an additional thirty minutes at 150°C, a vacuum was drawn to remove unreacted styrene and other lights. During the vacuum stripping, 8.5 ml of unreacted styrene and other lights were recovered in an overhead cold trap. After thirty minutes of vacuum stripping at 150°C, bisphenol A (162.8 g, 0.71 mole) was charged to the stirred solution. Three minutes later, the reactor temperature had dropped to 138°C with complete dissolution of the bisphenol A in the resin solution. An advancement catalyst (ethyltriphenylphosphonium acetate-acetic acid complex, 1 g, 70% in methanol) was added to the reactor which was then reheated to 150°C. Three minutes later, the reactor reached 150°C and heating was discontinued. A maximum exotherm of 173°C resulted six minutes later. The reactor was heated to 175°C where it

was held for one hour after the maximum exotherm. The polystyrene modified epoxy resin product was recovered at room temperature (25°C) as a pale-yellow colored, slightly opaque, brittle solid with a final epoxide equivalent weight of 708.

EXAMPLE 6

Cone and Plate Viscosity Determination of Powder Coating Resins

Cone and plate viscosity measurements were determined for the resins of Examples 1-5 and Comparative Experiment A utilizing an I.C.I. Cone and Plate Viscometer, Research Equipment (London) Ltd.

Viscosity measurements were taken at 150°C and 175°C. A 0.4 g sample of each solid resin was placed on the heated plate and evaluated with a size 100 cone. The Cone and Plate viscosity values for Examples 1-5 and Comparative Experiment A are reported in Table I.

EXAMPLE 7

Kinematic Viscosity Determination of Powder Coating Resins

Kinematic viscosity measurements were determined for the resin of Example 1 and a standard low gloss polystyrene modified epoxy resin prepared using the method of Comparative Experiment A utilizing Cannon-Fenske Routine Viscometers size 700.

Viscosity measurements were taken at 150°C. An eleven gram sample of each solid resin was placed into a size 700 viscometer. The viscometer was then inserted in to a 150°C constant temperature oil bath.

The viscometer was allowed to equilibrate at 150°C for at least sixty minutes prior to viscosity measurement. The molten resin was free of all air bubbles prior to viscosity measurement. The Kinematic viscosity values for Example 1 and the low gloss polystyrene modified epoxy resin prepared using the method of Comparative Experiment A are reported in Table IV.

EXAMPLE 8

Preparation of Powder Coating Formulations

Portions of each of the epoxy resins from Examples 1-5 and Comparative Experiment A were coarse ground in a Wellex Grinder. Formulation weights as per Table II, of the epoxy resin, dicyandiamide, a mixture of 17% by weight 2-methylimidazole and 83% by weight dicyandiamide, Modaflow II, and filler, were placed in a plastic bag, sealed and dry mixed to a homogeneous blend. The dry-mixed formulations were then extruded in a Buss-Condux PLK 46 single screw extruder (equipped with a 46 mm diameter kneader screw operated at 120 rpm). For the resin formulations of Examples 1-5, Zone 1 was set at 50°C and Zone 2 was set at 100°C; for the resin formulation of Comparative Experiment A, Zone 1 was set at 60°C and Zone 2 was set at 100°C. The extrudate was passed through BCI Chill Rolls (6 1/2" (165.1 mm) diameter, cooled and crushed). The crushed extrudate was then coarse ground in a Brinkmann Centrifugal Grinding Mill utilizing the 12-tooth attachment and then fine ground using the 12-tooth attachment and a 0.75 mm screen over the grinding teeth; dry ice was utilized as a coolant to minimize sintering and resin melt in the fine grinding step. The finely ground extrudate was sieved through No. 140 (150 mesh, 106 µm)

standard test sieves (wire cloth). The -150 mesh powder coating formulations were applied via electrostatic spray with a Gema Ag Type 710 Laboratory Unit (set at 60-70 kV) on to 4" x 12" x 20 gauge (101.6 mm x 304.8 mm x 0.529 mm) cold rolled steel, clean treatment Parker test panels (Parker Division, Hooker Chemicals and Plastics Corporation). The electrostatically coated panels were set in a Gallenkamp convection-type oven and cured at 180°C (356°F) for twenty minutes. After removal from the oven, the panels were cooled and evaluated via the following test methods: coating thickness was determined per ASTM D1186 by utilizing a Fischer Perma-Scope ES film thickness tester. Surface gloss was determined per ASTM D523 (DIN 67530) using a Mallinckrodt Multi Gloss gloss meter. Gardner forward and reverse impact strengths were determined 24 hours after spray-up as per ASTM D2794 using a Gardner "Coverall" Bend and Impact Tester, 46 inch (1.17 m) tube length, 0-160 in.-lb. tester, with a four pound (1.81 kg), one-half inch (12.7 mm) diameter cone. Visualization of any surface cracks at the impact sites was facilitated by application of an acidified copper sulfate (CuSO_4) solution for a period of 15 minutes. Impact areas were observed for copper deposits or iron-rust stains immediately after exposure to the copper sulfate solution.

Film thickness, surface gloss and Gardner impact strength values of the cured powder coatings prepared using the epoxy resins of Examples 1-5 and Comparative Experiment A are reported in Table II.

COMPARATIVE EXPERIMENT B

Preparation of Powder Coating Formulation Using Epoxy Resin Standard and 45 pbw TiO₂

5 A commercial grade (D.E.R.[®] 662UH, The Dow Chemical Company) diglycidyl ether of bisphenol A having an epoxide equivalent weight (EEW) of 714 was used to prepare a powder coating formulation using the method of Example 8 substituting titanium dioxide (TiO₂, DuPont R-900) for the BaSO₄ at 45 percent by 10 weight of the total formulation. Cured powder coated panels were prepared and tested using the methods of Example 8. The results are reported in Table III.

COMPARATIVE EXPERIMENT C

15 Preparation of Powder Coating Formulation Using Standard Low Gloss Epoxy Resin and 45 pbw TiO₂

20 A portion of resin from Comparative Experiment A was used to prepare a powder coating formulation using the method of Example 8 substituting titanium dioxide (TiO₂, DuPont R-900) for the BaSO₄ at 45 percent by weight of the total formulation. Cured powder coated steel panels were prepared and tested using the 25 methods of Example 8. The results are reported in Table III.

EXAMPLE 9

30 Preparation of Powder Coating Formulation Using 45 pbw TiO₂

35 A portion of resin from Example 5 was used to prepare a powder coating formulation using the method of Example 8 substituting titanium dioxide (TiO₂, DuPont R-900) for the BaSO₄ at 45 percent by weight of

the total formulation. Cured powder coated steel panels were prepared and tested using the methods of Example 8. The results are reported in Table III.

5

TABLE I

CONE AND PLATE VISCOSITY
cks (m^2/s)

10	<u>DESIGNATION OF EPOXY RESIN</u>	<u>150°C</u>	<u>175°C</u>	<u>EEW</u>
	Example 1	2400 (0.0024)	850 (0.00085)	721
	Example 2	3200 (0.0032)	1000 (0.001)	737
15	Example 3	3000 (0.003)	1000 (0.001)	733
	Example 4	3400 (0.0034)	900 (0.0009)	738
20	Example 5	2050 (0.00205)	750 (0.00075)	740
	Comparative Experiment A	*	1200 (0.0012)	708
	Comparative Experiment B	3500 (0.0035)	900 (0.00099)	714
25				

*An accurate viscosity reading could not be taken due to sample climbing viscosity cone.

30

35

TABLE II

Epoxy Resin Employed	Amount of Epoxy Resin grams	Curing Agent			Modaflow II ³ grams	Filler Type: grams	Filler Thickness mils (mm)	Gloss Degrees, Percent	Gardner Impact Forward/ Reverse in.-lb. (J)	Coating Appearance
		A ¹ grams	B ² grams							
Example 1	500	9.58	6.0		13.32	BaSO ₄ ; 150	1.2-2.2 (0.03-0.06)	20, 60, 85,	5.0 160/160 (18.08/18.08)	Flawless
Example 2	500	9.27	6.0		13.30	BaSO ₄ ; 150	1.3-1.7 (0.03-0.04)	20, 60, 85,	4.7 160/160 (18.08/18.08)	Flawless
Example 3	500	9.34	6.0		13.30	BaSO ₄ ; 150	1.2-2.0 (0.03-0.05)	20, 60, 85,	5.0 160/160 (15.82/13.56)	Flawless
Example 4	600	11.09	7.2		15.96	BaSO ₄ ; 180	1.2-2.2 (0.03-0.06)	20, 60, 85,	2.8 160/160 (18.08/18.08)	A
Example 5	500	9.21	6.0		13.30	BaSO ₄ ; 150	1.4-2.4 (0.04-0.06)	20, 60, 85,	4.7 160/160 (18.08/18.08)	Flawless
Comparative Experiment A	500	9.86	6.0		13.32	BaSO ₄ ; 150	1.4-2.2 (0.04-0.06)	20, 60, 85,	4.3 160/160 (18.08/18.08)	A

A = slight to barely perceptible texturing of surface



TABLE III

Epoxy Resin Employed	Amount of Epoxy Resin grams	Curing Agent		Modaflow II ³ grams	Filler Type: grams	Filler Thickness mils (mm)	Gloss Degrees, Percent	Gardner Impact Forward/ Reverse in.-lb. (J)	Coating Appearance
		A ¹ grams	B ² grams						
Example 5	500	9.21	6.0	18.72	TiO ₂ ; 421	1.3-2.1 (0.03-0.05)	20, 60, 85,	5.9 31.1 100/80 (11.3/9.04)	B,C
Comparative Experiment A	400	8.05	4.8	15.01	TiO ₂ ; 337.86	1.4-2.3 (0.04-0.06)	20, 60, 85,	1.6 7.4 100/40 (11.3/4.52)	B,C
Comparative Experiment B	400	7.78	4.8	15.00	TiO ₂ ; 337.56	1.3-2.0 (0.03-0.05)	20, 60, 85,	25.6 68.6 160/100 (18.08/11.3)	B

¹Curing Agent A was dicyandiamide

²Curing Agent B was mixture of 83 percent by weight dicyandiamide and 17 percent by weight 2-methylimidazole

³A polyacrylate flow control agent (Monsanto)

B = pronounced texturing of surface (orange peel)

C = cracking type failure noted at Gardner impact sites

TABLE IV

5	EPOXY RESIN EMPLOYED	KINEMATIC VISCOSITY AT 150°C	EEW
		cks (m^2/s)	
	Example 1	4541 (0.004541)	721
	Comparative Experiment A	7638 (0.007638)	702

10

EXAMPLE 10 - Inclined Plate Flow Test

15 Inclined plate flow was determined for the
resins of Example 1, Example 4, Example 5 and Compara-
20 tive Experiment A using the formulations prepared in
Example 8 and a herein described modification of
standard method ASTM D3451, Section 17. Samples (0.75
gram) of each resin were pressed in to 0.25" (6 mm)
25 thick by 0.50" (12.7 mm) diameter pellets utilizing a
Parr 2811 Pellet Press, stainless steel die (Parr
Instrument Co., Moline, Illinois). Three pellets of
each Example resin were prepared, six pellets of
Comparative Experiment A were prepared and used as
controls.

30 A constant temperature forced air electric oven
(Gruenberg Oven Co., Inc.) was set at 350°F. The inner
oven contained a metal plate rack assembly 5.7" (14.5
cm) x 6" (15.2 cm) which was capable of being
maintained in either a horizontal position or at a 65
degree angle by means of an exterior lever. A 4" x 12"
x 20 gauge (101.6 mm x 304.8 mm x 0.529 mm) polished,
35 S412, Q-panel (~~The Q-Panel Co., Cleveland, Ohio~~) was
cut in half (4" x 6" x 20 gauge (101.6 mm x 152.4 mm x

0.529 mm)) and placed in the preheated oven on the horizontal rack assembly. The panel was allowed to equilibrate for ten minutes at 350°F. After the panel had equilibrated at 350°F for ten minutes, three pellets one from Example 1, one from Example 4 and one from Comparative Experiment A were placed across the top of the preheated panel. The oven door was quickly closed, 30 seconds later the rack assembly holding the heated panel and pellets was tilted to 65 degrees from the horizontal via the exterior lever without opening the oven door. The rack, panel and pellets were allowed to remain in this position for 15 minutes. After 15 minutes, the panel was removed from the oven and allowed to cool to room temperature (25°C). The maximum flow, in millimeters, from the original diameter was measured. The same procedure was repeated for Example 5 and Comparative Experiment A. A series of three inclined plate flow tests was evaluated for each resin and the average millimeter flow value is reported in Table V.

TABLE V

25	<u>DESIGNATION OF EPOXY RESIN</u>	<u>AVERAGE INCLINED PLATE FLOW/STANDARD DEVIATION (mm)</u>
	Example 1	97.3/±1.8
	Example 4	69.8/±4.5
	Example 5	95.6/±4.7
30	Comparative Experiment A	57.1/±1.3

The embodiments of the invention in which an exclusive property or privilege is claimed are defined as follows:

1. An advanced epoxy resin composition which comprises the product resulting from

5 (I) polymerizing in the presence of a catalytic quantity of at least one suitable polymerization catalyst

(A) the reaction product of

(1) a mixture comprising

10 (a) from 80 to 99 percent by weight of at least one diglycidyl ether of a dihydric phenol having an epoxide equivalent weight of from 111 to 350 with the epoxide equivalent weight being
15 calculated on the basis that there are no substituent groups attached to the aromatic ring other than hydrogen and the
20 ether oxygen atoms even though the aromatic rings may in fact be substituted by groups other than hydrogen;

25 (b) from 1 to 20 percent by weight of at least one diglycidyl ether

of a mono-, di- or
polyetherdiol; with

- 5 (2) at least one compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group in an amount of from 0.001 to 0.05 equivalent per epoxide equivalent contained in component (A-1); with

10 (B) a monomer mixture comprising

- (1) at least one vinyl aromatic monomer in an amount of from 31 to 60 percent by weight of the combined weight of components (A), (B) and (C);
- 15 (2) at least one compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group in an amount of from 0.001 to 0.05 equivalent per epoxide equivalent contained in component (A-1); and optionally
- 20 (3) one or more hydroxyalkyl acrylate, hydroxyalkyl methacrylate, alkyl acrylate or alkyl methacrylate in an amount of from zero to 15 percent by weight of the combined weight of components (B-1) and (B-3); and
- 25 30

(II) advancing in the presence of a catalytic quantity of at least one suitable advancement catalyst, the product resulting from (I) with

- 35 (C) at least one dihydric phenol in an amount of from 0.125 to 0.8 hydroxyl equivalent

per epoxide equivalent in component (A-1).

2. An advanced epoxy resin of Claim 1 wherein
(i) component (A-1-a) is an epoxy resin or
mixture of epoxy resins represented by
the following formulas (I) or (II)

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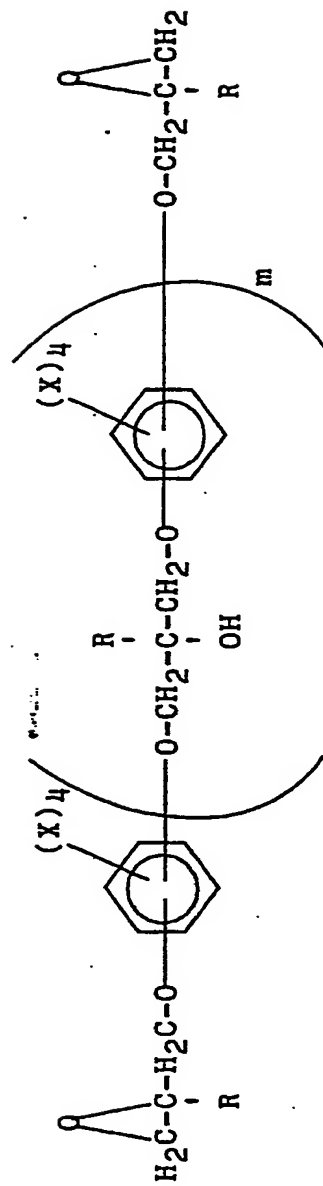
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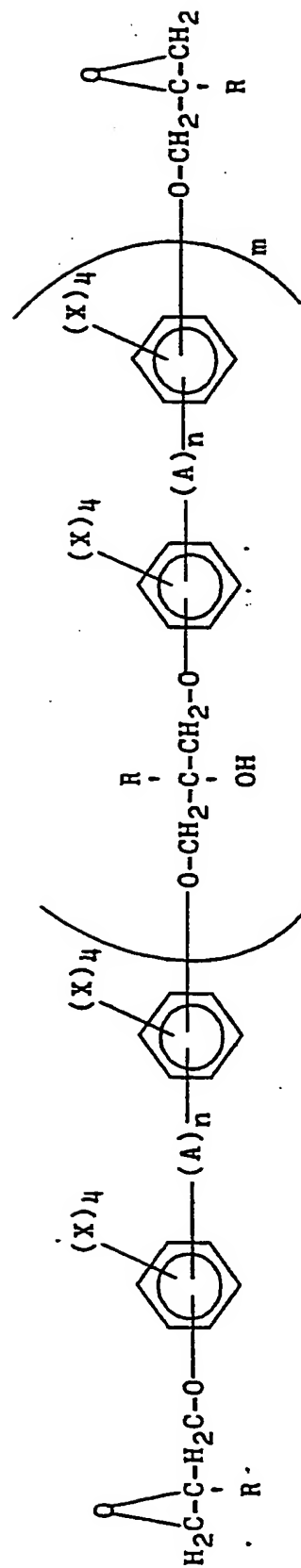
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(I)



(II)



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11

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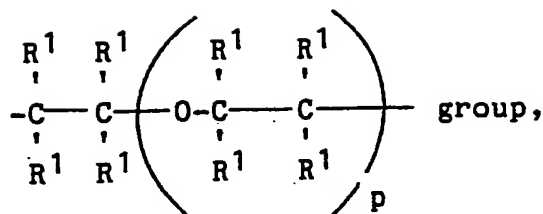
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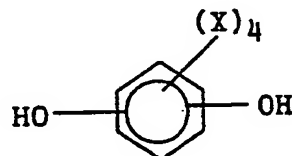
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R^1 is independently hydrogen or a hydrocarbyl group having from 1 to 6 carbon atoms, and p has a value from 1 to 100; m and R are as hereinbefore defined; and
(iii) component (C) is one or more dihydric phenols represented by the following formulas (VI) or (VII)

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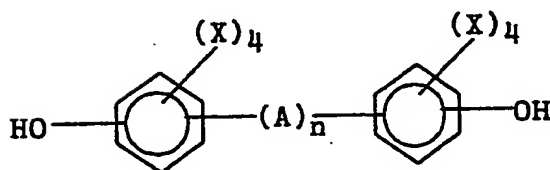
(VI)



25

30

(VII)



wherein X , A and n are as hereinbefore defined.

35

3. An advanced epoxy resin composition of Claim 2 wherein

- 5 (i) component (A-1-a) is a diglycidyl ether of bisphenol A, a diglycidyl ether of tetrabromobisphenol A, a diglycidyl ether of bisphenol F, or any combination thereof;
- 10 (ii) component (A-1-b) is a diglycidyl ether of a polyoxypropylene glycol, a diglycidyl ether of dipropylene glycol, a diglycidyl ether of a polyoxyethylene glycol, or any combination thereof;
- 15 (iii) component (A-2) is methacrylic acid, acrylic acid, or any combination thereof;
- (iv) component (B-1) is styrene, methylstyrene, or any combination thereof;
- 20 (v) component (B-2) is methacrylic acid, acrylic acid, or any combination thereof;
- 25 (vi) component (B-3) is hydroxyethyl acrylate, hydroxypropyl acrylate, hydroxyethyl methacrylate, hydroxypropyl methacrylate, lauryl methacrylate, or any combination thereof; and
- 30 (vii) component (C) is bisphenol A, tetrabromobisphenol A, bisphenol F, or any combination thereof.

4. An advanced epoxy resin composition which comprises the product resulting from

- 35 (I) polymerizing in the presence of a catalytic quantity of ~~at least one suitable polymerization~~ catalyst

(A) the reaction product of

(1) at least one diglycidyl ether of a dihydric phenol having an epoxide equivalent weight of from 111 to 350 with the epoxide equivalent weight being calculated on the basis that there are no substituent groups attached to the aromatic ring other than hydrogen and the ether oxygen atoms even though the aromatic rings may in fact be substituted by groups other than hydrogen;

(2) at least one compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group in an amount of from 0.001 to 0.05 equivalent per epoxide equivalent contained in component (A-1); with

(B) a monomer mixture comprising

(1) at least one vinyl aromatic monomer in an amount of from 31 to 60 percent by weight of the combined weight of components (A), (B) and (C);

(2) at least one compound containing both a group reactive with an epoxide group and a polymerizable ethylenically unsaturated group in an amount of from 0.001 to 0.05 equivalent per epoxide equivalent contained in component (A-1); and optionally

- 5 (3) one or more hydroxyalkyl acrylate,
hydroxyalkyl methacrylate, alkyl
acrylate or alkyl methacrylate in an
amount of from zero to 15 percent by
weight of the combined weight of
components (B-1) and (B-3); and
- 10 (II) advancing in the presence of a catalytic
quantity of at least one suitable
advancement catalyst, the product
resulting from (I) with a mixture
comprising
- 15 (C) at least one dihydric phenol in an
amount of from 0.125 to 0.8 hydroxyl
equivalent per epoxide equivalent in
component (A-1) and
- 20 (D) at least one diglycidyl ether of a mono-
, di- or polyetherdiol or at least one
phenolic capped adduct of a diglycidyl
ether of a mono-, di- or polyetherdiol
or combination thereof wherein the
25 amount of the diglycidyl ether or the
amount of the diglycidyl ether contained
in the capped diglycidyl ether is from 1
to 20 percent by weight of components
(I-A-1) and (II-D).
- 30
5. An advanced epoxy resin of Claim 4 wherein
- 35 (i) component (A-1-a) is an epoxy resin or
mixture of epoxy resins represented by the
following formulas (I) or (II)



wherein each A is independently a divalent hydrocarbon group having from one to 10 carbon atoms,
-O-, -S-, -S-S-,

5 0 0 0
 " " "

-S-, -S-, -C-; each X is independently

"

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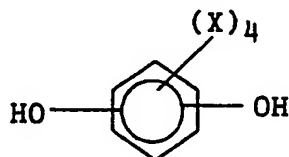
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hydrogen, bromine, chlorine, or a hydro-
carbonyl or hydrocarboxy group having from
1 to 10 carbon atoms; each R is
independently hydrogen or a methyl group;
15 m has a value from zero to 5; and n has a
value of zero or 1;

(ii) component (C) is one or more dihydric
phenols represented by the following
formulas (VI) or (VII)

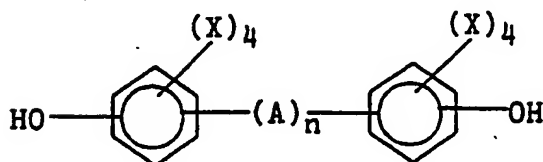
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(VI)



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(VII)



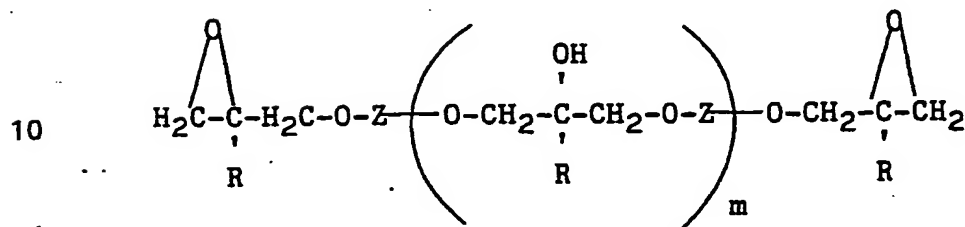
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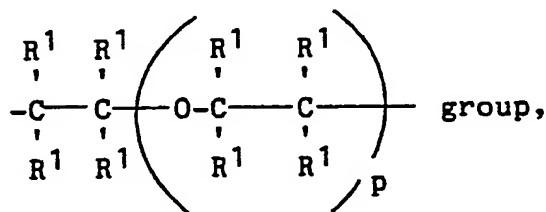
wherein X, A and n are as hereinbefore
defined; and

- (iii) component (D) is an epoxy resin or mixture of epoxy resins represented by the following formula (III)

(III)



wherein Z is a hydrocarbyl group containing from 1 to 12 carbon atoms or a



wherein each

R¹ is independently hydrogen or a hydrocarbyl group having from 1 to 6 carbon atoms, and p has a value from 1 to 100; m and R are as hereinbefore defined.

6. An advanced epoxy resin composition of Claim 5 wherein

- (i) component (A-1-a) is a diglycidyl ether of bisphenol A, a diglycidyl ether of tetrabromobisphenol A, a diglycidyl ether

of bisphenol F, or any combination thereof;

(ii) component (A-2) is methacrylic acid, acrylic acid, or combination thereof;

5 (iii) component (B-1) is styrene, methylstyrene, or any combination thereof;

(iv) component (B-2) is methacrylic acid, acrylic acid, or any combination thereof;

10 (v) component (B-3) is hydroxyethyl acrylate, hydroxypropyl acrylate, hydroxyethyl methacrylate, hydroxypropyl methacrylate, lauryl methacrylate, or any combination thereof;

15 (vi) component (C) is bisphenol A, tetrabromobisphenol A, bisphenol F, or any combination thereof; and

20 (vii) component (D) is a diglycidyl ether of a polyoxypropylene glycol, a diglycidylether of dipropylene glycol, a diglycidyl ether of a polyoxyethylene glycol, or any combination thereof.

25 7. A thermosettable (curable) composition comprising a composition of Claim 1, 2 or 3 and a curing amount of a suitable curing agent therefor.

30 8. A thermosettable (curable) composition of Claim 7 which is a powder coating formulation containing as a curing agent therefor a biguanide and optionally containing one or more adjuvants selected from the group consisting of curing agent accelerator or catalyst, particulate filler, pigment, flow control additive, gloss control additive, texture control
35 additive, air release additive, slip aid and the like.

9. A thermosettable (curable) composition of Claim 8 wherein said curing agent is dicyandiamide accelerated with 2-methylimidazole.

5 10. A thermosettable (curable) composition comprising a composition of Claim 4, 5, or 6 and a curing amount of a suitable curing agent therefor.

10 11. A thermosettable (curable) composition of Claim 10 which is a powder coating formulation containing as a curing agent therefor a biguanide and optionally containing one or more adjuvants selected from the group consisting of curing agent accelerator or catalyst, particulate filler, pigment, flow control
15 additive, gloss control additive, texture control additive, air release additive, slip aid and the like.

20 12. A thermosettable (curable) composition of Claim 11 wherein said curing agent is dicyandiamide accelerated with 2-methylimidazole.

13. A product resulting from curing the composition of Claim 7.

25 14. A product resulting from curing the composition of Claim 8.

15. A product resulting from curing the composition of Claim 9.

30 16. A product resulting from curing the composition of Claim 10.

35 17. A product resulting from curing the composition of Claim 11.

18. A product resulting from curing the composition of Claim 12.

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